

**ELECTROSTATIC PARTICLE CHARGER,
ELECTROSTATIC SEPARATION SYSTEM,
AND RELATED METHODS**

This application claims the benefit of U.S. Provisional Patent Application Ser. No. 60/477,443, the disclosure of which is incorporated herein by reference.

Technical Field

The present invention relates to the material separation art and, more particularly, to an improved particle charger or charging device, an improved separator, and related methods for electrostatically separating two
5 species of particles from a particle mixture.

Background of the Invention

“Dry” triboelectrostatic separation is widely used as an effective technique for separating different particulate solid components (“particles”) from a physical mixture entrained or carried in a driving fluid, such as air. Typical applications include the beneficiation of minerals, purification of
10 foods, the recovery of valuable components from waste, and the sizing of particles in a particle mixture. This technology has gained widespread acceptance as providing a low cost, environmentally friendly technique, since
15 it requires no chemicals or water and thus eliminates costly downstream dewatering and slime disposal applications required in wet separation processes.

Typically, electrostatic separation relies on the surface physical

properties of the different particles and controlled flow conditions to effect beneficiation in an efficient and effective manner. Specifically, when two species of particles with different work functions contact one another, a charge transfer between the contact area results, such that one species may carry a positive charge and the other a negative charge (known as "contact charging"). This differential charge may also be achieved by "friction charging," which results when the particles are forced to slide along or rub against a solid surface. The combined effects of these charges are together known as "triboelectrostatic charging" or "tribocharging" for short, and are together considered to play a key role in achieving particle separation.

Figure 1 schematically illustrates a typical prior art triboelectrostatic separator S. The particles P in the mixture are fed into the separator S from a bin B, and are charged to a bipolar state in a metal tube T, mainly by friction charging. The particles P then pass through an electric field F, such that the species of particles having a particular charge is drawn from the mixture toward a corresponding electrode E_1 , E_2 . However, as a result of the inefficient charging resulting from the fact that not all particles make contact with the sidewalls of the tube T, weakly charged or charge-neutral particles may not be attracted and consequently simply pass through the separator S unaffected by the electric field F. While these "middling" particles (not shown in Figure 1) may be separated during a second pass, this obviously decreases the efficiency of the separation operation. Increasing the feed rate of the particles P may allow for more passes in a shorter period, but a concomitant decrease in the separation efficiency per pass results because of the shorter residence time of the particles in the electric field F.

Accordingly, while the typical prior art separator S is effective for separating two particle species from a particle mixture, it should be appreciated that further improvements in separation effectiveness and

operational efficiency are still possible. More specifically, a need exists for devices and methods that enhance the charging on the particles as well as the downstream separation to improve efficiency and potentially reduce the need for the number of passes required.

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Summary of the Invention

In accordance with a first aspect of the invention, an apparatus for intended use in charging particles in a system for separating particles from a fluid flow is disclosed. The apparatus comprises: (1) a chamber including an
10 inlet for receiving the particles and an outlet for discharging the particles; and (2) a rotor rotatably mounted in the chamber. The rotor has a generally non-permeable outer surface for contacting and assisting in charging the particles.

In one particular embodiment, the rotor is circular, polygonal, or gear-shaped in cross-section, and the chamber is generally cylindrical.
15 Preferably, the outlet of the chamber is positioned below and generally opposite the inlet. A partition may also project into the chamber adjacent the rotor. Preferably, the partition is adjustable to vary the distance between an end of the partition and the rotor. Additionally, a motor is provided for rotating the rotor. The motor may rotate the rotor at a rotational speed of up
20 to 10,000 revolutions per minute.

In the same or another embodiment, an electric field is provided in the chamber. Preferably, the electric field is created by a variable voltage source having a first lead connected to the rotor and a second lead connected
25 to a wall of the chamber. The electric field helps to enhance the charging of certain types of particles.

In accordance with a second aspect of the invention, an apparatus for intended use in separating particles of a mixture is disclosed. The apparatus comprises a body including an inlet for receiving the electrically

charged particles to be separated, a separation chamber, a first electrode for attracting particles having a first selected charge, and a second electrode for attracting particles having a second selected charge. The first and second electrodes are grid electrodes having a plurality of elongated fingers extending
5 along the separation chamber spaced apart from the body. A flow straightener positioned in or adjacent to the inlet receives and straightens a co-flow of fluid, such as a gas, passing over and between the fingers of the grid electrodes for carrying or sweeping away the particles.

In one embodiment of the separation apparatus, a variable
10 voltage source applies a positive voltage potential to the first electrode and a negative voltage potential to the second electrode. Preferably, the fingers on each electrode are connected to a common header.

In accordance with a third aspect of the invention, a method of separating particles from a particle mixture is disclosed. The method
15 comprises actuating a rotor to create a differential charge on the two or more constituent species of particles in the mixture and separating the differentially charged particles into the two or more constituent species at a location downstream of the chamber. Preferably, the actuating step is accomplished by rotating the rotor at a speed of at least 1,200 revolutions per minute.

In accordance with a fourth aspect of the invention, a method for separating electrostatically charged particles from a mixture is disclosed. The method comprises introducing the charged particles to a separation chamber including a positive grid electrode for attracting negatively charged particles and a negative grid electrode for attracting positively charged particles; and
20 sweeping away corresponding particles from the grid electrodes using a
25 straightened co-flow of a fluid, such as a gas. The step of actuating a rotor in a mixing chamber upstream of the separation chamber to enhance the charge on the particles in the mixture may also be performed.

Brief Description of the Drawing Figures

Figure 1 schematically illustrates a prior art separation system;

Figure 2 is a partially cross-sectional, perspective view of one embodiment of the charger forming one aspect of the invention;

5 Figure 3 is a graph illustrating the enhanced particle charging achieved when an electric field is applied to the charger;

Figures 4a-4c show exemplary shapes of rotors;

Figure 5 is a partially cross-sectional, perspective view of one embodiment of the separator forming another aspect of the invention;

10 Figures 6a and 6b are schematic side views of the separator of Figure 5 in operation; and

Figure 7 illustrates an experimental set-up using the charger of Figure 2 and the separator of Figure 5.

15 **Detailed Description of the Invention**

With reference to the partially schematic, cross-sectional side view of Figure 2, and in accordance with a first aspect of the invention, an improved particle charging device or charger 10 is disclosed. The charger 10 includes a generally rectangular, elongated inlet 12 for receiving a feed stream
20 FS, which may include a mixture comprised of at least two species of particles to be differentially charged. Particles in the feedstream (which includes at least a small amount of a driving fluid, such as air) passing through the distributor are introduced to the inlet 12 and enter a charging chamber 14 forming part of the charger 10.

25 The charging chamber 14 is formed between the inner surface of an outer wall 16 and the outer surface of a charging roller or rotor 18 mounted to rotate about an axis of rotation X, and thus creates an annular space for receiving the particle mixture. The roller or rotor 18 is provided with a

generally continuous, non-permeable outer surface for contacting and frictionally charging the particles in the mixture (which typically have a size ranging from 2-3 millimeters or less).

5 An outlet 20 is defined in the outer wall 16 of the charger 10 generally opposite the inlet 12. The outlet 20 may be in direct or indirect communication with a downstream separator or like device for effecting further processing of the particle mixture. A plastic adaptor 22 may also be connected to the outlet 20 for receiving and containing the particle mixture as it transitions to the downstream separator S. To increase the throughput
10 without compromising efficiency, the charger 10 and all components forming it are elongated in a direction aligned with the axis of rotation of the rotor 18 (which is shown as being hollow and having a center support shaft (not numbered) in operative engagement at one end with a motor M).

In one possible mode of operation, the rotor 18 is rotated at a
15 selected rotational speed (e.g., up to 10,000 rpm, and more preferably between 1,200 and 8,000 rpm) by the motor M (which may be a variable speed electric motor). Particles encountering the rotor 18 upon passing through the inlet 12 become agitated and charged by both friction and contact charging. More particularly, the dynamic agitation of the mixture created by the rotation of the
20 rotor 18 increases the incidence of both: (1) particle-particle contact, thus creating contact charging; and (2) particle-wall contact (either the outer wall 16 or with the surface of the rotor 18), thus creating friction charging. In other words, the particles in the mixture will have multiple areas of contact, both with the rotor 18 and the other particles, due to the fast rotation and agitation
25 of the particles created thereby. As a result of using this "rotary charger," a much higher charge density on the surface of the particles results, and the incidence of weakly or neutrally charged particles passing through the outlet 20 is reduced.

When the particles passing through the charger 10 are fed to a downstream separator S, separation efficiency is increased (possibly by as much as 40%) and the need for multiple passes to effect separation may be eliminated. The active charging provided by the charger 10 also allows for a much higher throughput without reducing the separation efficiency, as compared to the passive charging afforded by the tube-type of arrangement shown in Figure 1. The charger 10 also helps to ensure that all particles are charged, not just a mono-layer of particles at the surface of the mixture (as is the case of a corona charger).

The charger 10 may also operate in a continuous fashion such that particles fed through the inlet are constantly being charged and discharged through the outlet for downstream separation. However, the provision of a closure or door adjacent the outlet 20 is a possibility, including in the case where the operation of the charger is separate from the downstream operation. In other words, the charging may be completed apart from the separation, the two may occur simultaneously on the same batch of the particle mixture, or the two may occur simultaneously on two different batches of the particle mixture.

Figure 2 also illustrates that a partition 24 may also be provided for selective insertion into the chamber 14 to perform the dual function of preventing the particle mixture from prematurely entering the outlet 20 in one direction and guiding the particle mixture to the outlet in the other. The partition 24 may pass through an opening in the outer wall 16, preferably adjacent to the opening defined by the outlet 20 through which the particle mixture exits the chamber 14, and its inner end extends to a point closely adjacent to the outer surface of the rotor 18. This inner end of the partition 24 may have an upper face matching the contour of the rotor (e.g., an arcuate face, in the case where the rotor is cylindrical)). The partition 24 may be mounted directly to the wall defining the outlet 20 using a fastener (Figure 2), and may

optionally be mounted to permit selective adjustment of the inner end toward or away from the rotor 18.

When the rotor 18 rotates in the clockwise direction as viewed in Figure 2 (note action arrow A), the partition 24 is thus positioned downstream of the outlet 20 in the angular sense. In this position, it serves to prevent or block particles from simply falling through the outlet 20 without making contact with the surface of the rotor 18 or the inside surface of the outer wall 16. The partition 24 so positioned also prevents lighter particles from becoming permanently suspended in the fluid flow surrounding the rotor 18 during rotation, since it contacts and forces the particles into the outlet 20 and toward the downstream separator. As should be appreciated, when the direction of rotation is reversed, the position of the partition 24 relative to the outlet 20 may be likewise reversed to accomplish the intended blocking and guiding functions.

Selective charging may further be enhanced by applying an electric field to the charger 10. Specifically, as shown in Figure 2, the leads of an external voltage source 26 are applied to the rotor 18 and the outer wall 16 of the chamber 14 to create an electric field therein. Using this externally applied voltage may allow for a certain charge density and polarity to be achieved on the particles. For example, as graphically illustrated in Figure 3, using a typical phosphate and quartz mixture, the phosphate may be charged over a range of 500×10^{-6} C/kg (from about positive 250×10^{-6} C/kg to negative 250×10^{-6} C/kg) while the quartz is always charged negatively when the external applied voltage is in the range of -9kV to +9kV. At zero voltage, both the phosphate and quartz are charged negatively. Therefore, separation of phosphate from quartz is more efficient if an external voltage is applied.

Although a generally cylindrical rotor 18 is shown in Figure 2, it should be appreciated that other shapes may be used (and that such shapes

may further enhance the charging of the particles). An example of a gear or sprocket-wheel type of rotor 18a with outwardly projecting "teeth" and corresponding surface indentations is shown in Figure 4a. An octagonal rotor 18b is shown in Figure 4b, which thus provides a plurality (eight) of distinct contact surfaces. Other polygonal shapes could also be used (e.g., a hexagon or dodecagon), with or without rounded corners at the intersections of the planar surfaces. Figure 4c also shows that a cylindrical separator 18c may be formed as a solid body, as compared to the hollow body 18 shown in Figure 2. In both cases, the outer surface of the rotor 18 remains non-permeable or continuous.

In accordance with another aspect of the invention, an improved separator 100 is also disclosed. The separator 100 includes a distributor 112 defining an inlet for receiving a feedstream of charged particles (which as should be appreciated may be delivered from the outlet 20 of the charger 10 described above or a different device, including the conventional tube T shown in Figure 1). The particles are delivered to a body 114 of the separator 100, which is generally rectangular and elongated. The driving fluid (gas) may be supplied by a driving fluid source, such as a forced draft fan (not shown), positioned upstream of the distributor 112. The fluid or gas is preferably ambient air, but other gases such as nitrogen, helium, argon, carbon dioxide, or combustion flue gas can be used at temperatures between approximately 25° C to 300° C.

In typical separators using plate-type electrodes (see Figure 1), the deposition of charged particles may reduce the separation efficiency and require deleterious periodic shutdowns for cleaning. In an effort to reduce or eliminate the need for such shutdowns, a pair of grid or grill electrodes 116, 118 are positioned in the body 114, spaced from the sidewalls thereof. Each grid electrode 116, 118 is comprised of a plurality of elongated, generally

parallel fingers 120 that extend into and define a separation chamber 122 within the body 114. The fingers 120 each emanate from a common header 124 and a similar footer (not shown) may also be provided to enhance the rigidity and stability of the electrodes 116, 118. The elongated, spaced nature
5 of the fingers 120 eliminates or substantially reduces the amount of particles deposited on the electrodes 116, 118, which obviates the above-mentioned problems.

Each electrode 116, 118 is connected to the lead of a variable voltage source 126 (such as along the header 124) to create an electric field in
10 the chamber 122 for separating the particles having a selected charge. A co-flow of gas devoid of particles may also be introduced from a separate source (not shown) for sweeping away the particles drawn towards the electrodes 116, 118. Preferably, flow straighteners 128 are provided to reduce the turbulence and form a smooth co-flow of gas generally parallel to the feedstream FS upon
15 entering the separation chamber 122. The flow straighteners 128 may be in the form of tubes having aspect ratios, i.e., the ratio of length to diameter, of greater than 20:1, but other types of straighteners (such as vanes) may also be used.

Figures 6a and 6b schematically demonstrate a comparison
20 between the use of plate electrodes P (Figure 6a) and the grill or grid electrodes 116, 118 (Figure 6b). In Figure 6a, the deposit D of particles on the plate electrodes P is shown. Since the flow of both the feedstream FS of charged particles and the co-flow CF devoid of particles passes only over the opposed faces of the plate electrodes P, the particles drawn from the mixture
25 accumulate and form the deposits D. However, in the case of the grid electrodes 116, 118 (which are spaced from at least two adjacent sidewalls of the body 114, and preferably all four), the flows essentially surround the fingers, moving both over and between them. This helps to prevent the

particles from accumulating and forming the undesirable deposits that hamper efficient operation.

Figure 7 shows an experimental set-up built and used to demonstrate the effectiveness of the charger 10 and separator 100 disclosed herein when used in combination. As is known in the art, the system may include a splitter 140 downstream of the separator for dividing the flow into streams including the substantially separate and pure species of particles and cyclones 150 or other filtering devices for removing the particles from the streams once separated. Collection bins 160 may also be provided for collecting the first and second species of particles, as well as any "middlings" that result.

Experiments were conducted using the exemplary system 100 shown in Figure 7 in an effort to demonstrate the efficacy of the charger 10. Both one-stage and two-stage separation was employed. Using this set-up, the following sets of data were obtained using both one-stage and two-stage separation:

Example 1 - Fly ash separation result

Table 1. One-stage fly ash separation

#	Ash		Middling		Tailing	
	LOI, %	Yield	LOI, %	Yield	LOI, %	Yield
#1	0.75	44.57	1.92	43.30	4.31	12.13
#2	0.94	59.92	7.49	25.62	39.19	14.46
#3	1.19	33.87	2.82	38.09	15.98	28.04
#4	1.21	42.08	5.37	45.82	28.47	12.10
#5	3.64	43.41	14.93	39.22	41.44	17.37

Example 2 - Fly ash separation result

Table 2. Two-stage fly ash separation

Product	Σ Product	Product	Σ Product	Σ Ash
Ash, %	Ash, %	Yield, %	Yield, %	Recovery, %
0.23	0.23	36.01	36.01	38.85
1.04	0.48	15.58	51.59	55.52
3.58	0.72	4.48	56.06	60.18
6.04	1.31	6.90	62.96	67.19
7.12	1.67	4.20	67.16	71.41
9.64	3.11	14.85	82.01	85.92
13.03	3.73	5.45	87.45	91.04
27.13	4.50	2.98	90.43	93.38
30.84	5.94	5.24	95.67	97.30
42.38	7.52	4.34	100.00	100.00

Example 3 - Coal cleaning result

Table 3. Coal cleaning

Product	Σ Product	Product	Σ Product	Σ Combustible	Σ Ash
Ash, %	Ash, %	Yield, %	Yield, %	Recovery, %	Rejection, %
3.44	3.44	42.40	42.40	49.56	91.61
7.82	4.84	19.94	62.34	71.81	82.64
26.92	9.07	14.75	77.10	84.86	59.78
37.89	13.02	12.24	89.34	94.06	33.09
53.96	17.38	10.66	100.00	100.00	0.00

Table 3 shows the results of coal cleaning obtained by a two-stage closed circuit test. The raw coal ash content is about 17%. For the product with 9.07% ash, an 84.86% of combustible recovery can be achieved with an ash rejection of 59.78%.

Example 4 - Ground calcium carbonate separation result

Table 4. Separation results on ground calcium carbonate (GCC)

	Insoluble %	Σ Insoluble %	Yield %	Σ Yield %	Σ Recovery %
5					
One-Stage	0.50	0.50	40.70	40.70	41.99
	3.00	1.88	50.10	90.79	92.39
	20.30	3.58	9.21	100.00	100.00
			100.00		
10					
Two-Stage	0.10	0.10	14.61	14.61	15.11
	0.50	0.33	19.61	34.22	35.31
	0.50	0.39	19.20	53.42	55.09
	1.80	0.50	4.34	57.76	59.51
	2.50	1.16	28.31	86.07	88.09
	6.40	1.35	3.37	89.43	91.35
	12.50	1.83	3.99	93.42	94.96
	16.60	2.54	4.71	98.13	99.03
15	49.90	3.42	1.87	100.00	100.00
			100.00		

As shown in Table 4, efficient removal of silica from the ground calcium carbonate (GCC) was achieved with the triboelectrostatic separation technology. A two-stage separation produced better separation results than the one-stage separation. Based on the two-stage separation, approximately 34% of calcium carbonate can be recovered for a product with 0.3% insol; a 57% yield of calcium carbonate is expected for a product with 0.5% insoluble.

Example 5 - Phosphate separation result

Table 5. Two-stage separation on phosphate flotation feed

	P ₂ O ₅ %	Σ P ₂ O ₅	Yield %	Σ Yield	Σ P ₂ O ₅ Recovery %
30	36.64	36.64	5.15	5.15	32.35
	17.32	23.48	11.00	16.14	65.03
	14.42	21.48	4.55	20.70	76.29
	12.96	19.99	4.40	25.09	86.06
	3.21	13.82	14.58	39.67	94.08
	1.99	12.26	6.03	45.70	96.14
	1.01	10.47	8.64	54.34	97.64
35	0.36	7.23	25.68	80.02	99.23
	0.22	5.83	19.98	100.00	100.00

Two-stage separation was conducted on a phosphate sample (Table 5), which is the flotation feed. Two fractions containing less than 0.5% P_2O_5 with 45% yield exist. A concentrate with 36.64% P_2O_5 can be produced with 32.35% P_2O_5 recovery.

5 The foregoing descriptions of various embodiments of the invention are provided for purposes of illustration, and are not intended to be exhaustive or limiting. Modifications or variations are also possible in light of the above teachings. The embodiments described above were chosen to provide the best application to thereby enable one of ordinary skill in the art to
10 utilize the disclosed inventions in various embodiments and with various modifications as are suited to the particular use contemplated. All such modifications and variations are within the scope of the invention as determined by the appended claims when interpreted in accordance with the breadth to which they are fairly, legally and equitably entitled.